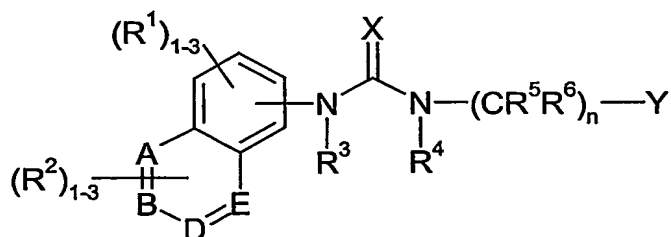


CLAIMS

1. A compound of formula (I):



(I)

5

wherein

A, B, D and E are each C or N with the proviso that one or more are N;

R¹ and R² are each independently hydrogen, halogen, hydroxy, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, haloC₁₋₆alkyl, hydroxyC₁₋₆alkyl, C₁₋₆alkoxy,

10 haloC₁₋₆alkoxy, hydroxyC₁₋₆alkoxy, C₃₋₇cycloalkyl, C₃₋₅cycloalkylC₁₋₄alkyl, NR⁷R⁸, carboxy, esterified carboxy, C₁₋₆alkyl substituted with a group selected from NR⁷R⁸, carboxy and esterified carboxy, or C₁₋₆alkoxy substituted with a group selected from NR⁷R⁸, carboxy and esterified carboxy;

R³ and R⁴ are each independently hydrogen, C₁₋₆alkyl, C₂₋₆alkenyl or C₂₋₆alkynyl;

15 R⁵ and R⁶ are, at each occurrence, independently hydrogen, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, C₁₋₆alkoxy, C₁₋₆acyloxy, carboxy, esterified carboxy, CONR⁷R⁸, SO₂R⁷, SO₂NR⁷R⁸, aryl, heteroaryl, heterocyclyl, or C₁₋₆alkyl substituted with a group selected from hydroxy, C₁₋₆alkoxy, C₁₋₆acyloxy, carboxy, esterified carboxy, NR⁷R⁸, CONR⁷R⁸, SR⁷, SO₂R⁷, SO₂NR⁷R⁸, aryl, heteroaryl and heterocyclyl;

20 or R⁵ and R⁶ and the carbon atom to which they are attached together form a carbocyclic ring of 3 to 6 carbon atoms;

R⁷ and R⁸ are, at each occurrence, independently hydrogen, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, C₃₋₇cycloalkyl or fluoroC₁₋₆alkyl;

or R⁷ and R⁸ and the nitrogen atom to which they are attached together form a

25 heteroaliphatic ring of 4 to 7 ring atoms, optionally substituted by one or two groups selected from hydroxy or C₁₋₄alkoxy, which ring may optionally contain as one of the said ring atoms an oxygen or a sulfur atom, a group S(O) or S(O)₂, or a second nitrogen atom which will be part of a NH or NR^a moiety where R^a is C₁₋₄alkyl optionally substituted by hydroxy or C₁₋₄alkoxy;

X is an oxygen or sulfur atom or the group =NCN;

Y is an aryl, heteroaryl, carbocyclyl or fused-carbocyclyl group; and

n is either zero or an integer from 1 to 3;

or a pharmaceutically acceptable salt, N-oxide or a prodrug thereof.

5

2. A compound according to claim 1 in which X is O.

3. A compound according to claim 1 or 2 in which R³ and R⁴ are hydrogen.

10 4. A compound according to claim 1, 2 or 3 in which B is nitrogen and A, D and E are carbon.

5. A compound according to any preceding claim in which Y is an aryl group selected from unsubstituted phenyl or naphthyl and phenyl or naphthyl
15 substituted by one or two substituents selected from halogen, C₁₋₄alkyl, C₁₋₄alkoxy, haloC₁₋₄alkyl, haloC₁₋₄alkoxy, phenyl, cyano, nitro, pyrazolyl, di(C₁₋₆alkyl)amino, phenoxy, -OCH₂O- and C₁₋₆alkylcarbonyl; or a heteroaryl group selected from pyridyl, thiazolyl, isoxazolyl, oxadiazolyl and pyrazolyl wherein each heteroaryl group is optionally substituted with one or two
20 substituents selected from C₁₋₄alkyl, C₁₋₄alkoxy, haloC₁₋₄alkyl, haloC₁₋₄alkoxy, unsubstituted heteroaryl or phenyl which may be substituted by C₁₋₆alkyl or halogen; or a carbocyclyl group which is a C₅₋₇cycloalkyl radical that is unsubstituted or substituted by a phenyl ring; or a fused-carbocyclyl group which is a C₅₋₇cycloalkyl radical that is fused to a phenyl ring.

25

6. A compound according to any preceding claim wherein R⁵ and R⁶ each independently represent a hydrogen atom or a C₁₋₄alkyl or phenyl group.

7. A pharmaceutical composition comprising a compound according to any
30 preceding claim or a pharmaceutically acceptable salt or N-oxide thereof.

8. A compound according to any one of claims 1 to 6 or a pharmaceutically acceptable salt or N-oxide thereof for use in a method of treatment of the human or animal body by therapy.

9. Use of a compound according to any one of claims 1 to 6 or a
pharmaceutically acceptable salt or N-oxide thereof for use in the manufacture of
a medicament for treating diseases and conditions in which pain and/or
5 inflammation predominates.
10. A method of treating a subject suffering from a disease or condition in
which pain and/or inflammation predominates which comprises administering to
that subject a therapeutically effective amount of a compound according to claim
10 1 or a pharmaceutically acceptable salt or N-oxide thereof.